

In the Claims:

Claims 1-70 (canceled).

71. (Previously presented) A method of inhibiting growth of a refractory tumor that has failed or been resistant to treatment with chemotherapy or radiation therapy comprising administering to a human, without concomitant chemotherapy or radiation therapy, an epidermal growth factor receptor (EGFR) antagonist that is an anti-EGFR antibody, or fragment thereof that retains the ability to bind to the EGFR, wherein administration is effective to inhibit growth of the refractory tumor.

72. (Previously presented) The method according to claim 71, wherein the refractory tumor overexpresses EGFR.

73. (Previously presented) The method according to claim 71, wherein the refractory tumor is a refractory tumor of the breast, heart, lung, small intestine, spleen, kidney, bladder, ovary, prostate, brain, pancreas, skin, bone, bone marrow, blood, thymus, uterus, testicles, cervix, or liver.

74. (Previously presented) The method according to claim 71, wherein the refractory tumor is a refractory tumor of the breast, heart, lung, small intestine, spleen, kidney, bladder, ovary, brain, pancreas, skin, bone, bone marrow, blood, thymus, uterus, testicles, cervix, or liver.

75. (Previously presented) The method according to claim 71, wherein the refractory tumor is a refractory squamous cell tumor.

76. (Previously presented) The method according to claim 71, wherein the EGFR antagonist is administered intravenously.

77. (Previously presented) The method according to claim 71, wherein the EGFR antagonist is administered at a dose of about 10 to about 500 mg/m² weekly.

78. (Previously presented) The method according to claim 71, wherein the EGFR antagonist inhibits stimulation of EGFR by its ligand.

79. (Previously presented) The method according to claim 78, wherein the EGFR antagonist inhibits binding of EGFR to its ligand.

80. (Previously presented) The method according to claim 78, wherein the EGFR antagonist binds EGFR externally.

81. (Previously presented) The method according to claim 78, wherein the EGFR antagonist inhibits EGFR phosphorylation.

82. (Previously presented) The method according to claim 78, wherein the EGFR antagonist inhibits EGFR tyrosine kinase activity.

83. (Previously presented) The method according to claim 71, wherein the anti-EGFR antibody comprises a constant region of a human antibody.

84. (Previously presented) The method according to claim 83, wherein the antibody is a chimeric antibody further comprising a variable region of a non-human animal antibody.

85. (Previously presented) The method of claim 84, wherein the non-human animal antibody is a mouse antibody.

86. (Currently amended) The method according to claim 83, wherein the antibody is a humanized antibody further comprising a variable region having all six complementarity-determining regions (CDRs) of a non-human animal antibody and a framework variable region of a human antibody.

87. (Previously presented) The method of claim 86, wherein the non-human animal antibody is a mouse antibody.

88. (canceled).

89. (Currently amended) The method according to claim 71, wherein the antibody is a human antibody having six complementarity-determining regions (CDRs).

90. (Previously Presented) The method according to claim 71, wherein the anti-EGFR antibody is administered at a dose sufficient to saturate EGFR.

91. (Currently amended) The method according to claim 71, wherein the method further comprises administering an adjuvant that is an immune system stimulator.

92. (Previously presented) The method of claim 71, wherein the EGFR antagonist comprises the anti-EGFR antibody.

93. (Previously presented) The method of claim 71, wherein the EGFR antagonist comprises the fragment of the anti-EGFR antibody.

94. (Previously presented) The method of claim 93, wherein the fragment comprises one or both Fab fragments or the F(ab')₂ fragment of the anti-EGFR antibody.

95. (Previously presented) The method of claim 93, wherein the fragment is a single chain antibody.

96. (Previously presented) The method of claim 71, wherein the refractory tumor is a carcinoma, glioma, sarcoma, adenocarcinoma, adenosarcoma or adenoma.

97. (Previously presented) A method of inhibiting growth of a refractory tumor that has failed or been resistant to treatment with chemotherapy or radiation therapy comprising administering to a human, without concomitant chemotherapy or radiation therapy, an epidermal growth factor receptor (EGFR) antagonist that is an anti-EGFR antibody, or fragment thereof that retains the ability to bind to the EGFR, wherein administration is effective to inhibit growth of the refractory tumor, wherein the refractory tumor is a refractory tumor of the colon.

98. (Previously presented) A method of inhibiting growth of a refractory tumor that has failed or been resistant to treatment with chemotherapy or radiation therapy comprising administering to a human, without concomitant chemotherapy or radiation therapy, an epidermal growth factor receptor (EGFR) antagonist that is an anti-EGFR antibody, or fragment thereof that retains the ability to bind to the EGFR, wherein administration is effective to inhibit growth of the refractory tumor, wherein the refractory tumor is a refractory tumor of the head and neck.